U.S. Application Serial No. 09/333,159

Attorney Docket No. MBI099-030RCEM

Marked Up Copy of Claims, as Amended in the Amendment Filed in Response to the Office Action Dated February 14, 2003

- 1. (Amended Four Times) An isolated nucleic acid molecule, or its complement, wherein the isolated nucleic acid i) encodes a polypeptide which exhibits lipase activity and ii) is selected from the group consisting of:
- a) a nucleic acid molecule having a nucleotide sequence which is at least 90% identical to the nucleotide sequence of SEQ ID NO: 45 or 46;
 - b) a nucleic acid molecule comprising a fragment of SEQ ID NO: 45 or 46;
- c) a nucleic acid molecule which encodes a polypeptide comprising the amino acid sequence encoded by SEQ ID NO: 46;
- d) a nucleic acid molecule which encodes a fragment of the amino acid sequence encoded by SEQ ID NO: 46; and
- e) a nucleic acid molecule which encodes a variant of the amino acid sequence encoded by SEQ ID NO: 46, wherein the nucleic acid molecule hybridizes over its full length in 6× sodium chloride/sodium citrate (SSC) at about 45°C, followed by one or more washes in 0.2× SSC, 0.1% SDS at 50°C with a portion of a nucleic acid molecule consisting of the nucleotide sequence of SEQ ID NO: 45 or 46.
- 39. (Twice Amended) The isolated nucleic acid molecule of claim 1, or its complement, wherein the molecule hybridizes over its full length in 6× SSC at about 45°C, followed by one or more washes in 0.2× SSC, 0.1% SDS at 50°C with a nucleic acid molecule consisting of the nucleotide sequence of SEQ ID NO: 45 or 46.

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- 33. (Twice Amended) The isolated nucleic acid molecule of claim 30, or its complement, wherein the <u>nucleic acid molecule encodes a polypeptide comprising a fragment</u> which comprises consecutive amino acid residues comprise an immunogenic portion of the protein having the amino acid sequence encoded by SEQ ID NO: 46.
- 32. (Thrice Amended) The isolated nucleic acid molecule of claim 1, or its complement, wherein the nucleic acid molecule encodes a variant of the amino acid sequence encoded by SEQ ID NO: 46, wherein the nucleic acid molecule hybridizes over its full length in 6× SSC at about 45°C, followed by one or more washes in 0.2× SSC, 0.1% SDS at 50°C with a nucleic acid molecule consisting of the nucleotide sequence of SEQ ID NO: 45 or 46.
- 37. (Thrice Amended) The method of claim 12, wherein the polypeptide is a variant of the polypeptide encoded by SEQ ID NO: 46, wherein the polypeptide is encoded by a nucleic acid molecule which hybridizes over its full length in 6× SSC at about 45°C, followed by one or more washes in 0.2× SSC, 0.1% SDS at 50°C with a nucleic acid molecule consisting of the nucleotide sequence of SEQ ID NO: 45 or 46, or a complement thereof.
- 43. (Amended) An isolated nucleic acid molecule, or its complement, wherein the isolated nucleic acid i) encodes an immunogenic portion of the protein having the amino acid sequence encode by SEQ ID NO: 46 and ii) is selected from the group consisting of:
- a) a nucleic acid molecule having a nucleotide sequence which is at least 90% identical to the nucleotide sequence of SEQ ID NO: 45 or 46;
 - b) a nucleic acid molecule comprising a fragment of SEQ ID NO: 45 or 46;
- c) a nucleic acid molecule which encodes a polypeptide comprising the amino acid sequence encoded by SEQ ID NO: 46;

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- d) a nucleic acid molecule which encodes a fragment of the amino acid sequence encoded by SEQ ID NO: 46; and
- e) a nucleic acid molecule which encodes a variant of the amino acid sequence encoded by SEQ ID NO: 46, wherein the nucleic acid molecule hybridizes over its full length in 6× sodium chloride/sodium citrate (SSC) at about 45°C, followed by one or more washes in 0.2× SSC, 0.1% SDS at 50°C with a portion of a nucleic acid molecule consisting of the nucleotide sequence of SEQ ID NO: 45 or 46.
- 44. (Amended) The isolated nucleic acid molecule of claim 43, or its complement, wherein the molecule hybridizes over its full length in 6× SSC at about 45°C, followed by one or more washes in 0.2× SSC, 0.1% SDS at 50°C with a nucleic acid molecule consisting of the nucleotide sequence of SEQ ID NO: 45 or 46.
- 54. (Amended) The isolated nucleic acid molecule of claim 43, or its complement, wherein the nucleic acid molecule encodes a variant of the amino acid sequence encoded by SEQ ID NO: 46, wherein the nucleic acid molecule hybridizes over its full length in 6× SSC at about 45°C, followed by one or more washes in 0.2× SSC, 0.1% SDS at 50°C with a nucleic acid molecule consisting of the nucleotide sequence of SEQ ID NO: 45 or 46.
- 65. (Twice Amended) The method of claim 62, wherein the <u>immunogenic</u> portion is from a polypeptide is a variant of the polypeptide encoded by SEQ ID NO: 46, wherein the polypeptide is encoded by a nucleic acid molecule which hybridizes <u>over its full</u> length in 6× SSC at about 45°C, followed by one or more washes in 0.2× SSC, 0.1% SDS at 50°C with a nucleic acid molecule consisting of the nucleotide sequence of SEQ ID NO: 45 or 46, or a complement thereof.

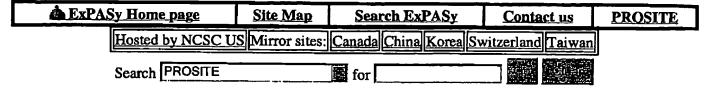
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General inform	ation about the entry
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Accession number	PS00120
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PROSITE documentation	PDOC00110
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Number of different Number of different Number of Number of Number of Precision Recall (true) Taxonomics Maximum	ther of hits in SWISS-PROT: 113 hits in 113 different sequences If hits on proteins that are known to belong to the set under consideration: 73 hits in 73 sequences If hits on proteins that could potentially belong to the set under consideration: 2 hits in 2 sequences If false hits (on unrelated proteins): 38 hits in 38 different sequences If known missed hits: 10 If partial sequences which belong to the set under consideration, but which are not hit by the profile because they are partial (fragment) sequences: 2 If true hits / (true hits + false positives)): 65.77 % If the hits / (true hits + false negatives)): 87.95 % If crange: Eukaryotes, Prokaryotes (Bacteria) If known number of repetitions of the pattern in a single protein: 1 If y'site in the pattern: 7,active_site
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Detailed view
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LEGAL DEPT.

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DR
DR
DR
          P40363, YJG8_YEAST, F; Q10687, YK79_MYCTU, F; Q03565, YKD7_CAEEL, F;
DR
          Q20076, YL4K_CAEEL, F; Q10508, YM23_MYCTU, F; Q10509, YM24_MYCTU, F; P40345, YN84_YEAST, F; P23974, YTXM_BACSU, F; 1RP1; 1BU8; 10IL; 2LIP; 3LIP; 1TAH; 1TGL; 3TGL; 4TGL; 5TGL; 1LGY; 1TIB;
DR
 DR
 3D
           1TIC; 1HPL; 1LPA; 1LPB; 1ETH; 1TIA;
 3D
          PDOC00110;
 DO
 {PDOC00110}
 {PS00120; LIPASE_SER}
 {BEGIN}
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Page 2 of 2

Triglyceride lipases (EC 3.1.1.3) [1] are lipolytic enzymes that hydrolyzes the ester bond of triglycerides. Lipases are widely distributed in animals, plants and prokaryotes. In higher vertebrates there are at least three tissue-specific isozymes: pancreatic, hepatic, and gastric/lingual. These three types of lipases are closely related to each other as well as to lipoprotein lipase (EC 3.1.1.34) [2], which hydrolyzes triglycerides of chylomicrons and very low density lipoproteins (VLDL).

The most conserved region in all these proteins is centered around a serine residue which has been shown [3] to participate, with an histidine and an aspartic acid residue, to a charge relay system. Such a region is also present in lipases of prokaryotic origin and in lecithin-cholesterol acyltransferase (EC 2.3.1.43) (LCAT) [4], which catalyzes fatty acid transfer between phosphatidylcholine and cholesterol. We have built a pattern from that region.

- -Consensus pattern: [LIV]-x-[LIVFY]-[LIVMST]-G-[HYWV]-S-x-G-[GSTAC]
 [S is the active site residue]
- -Sequences known to belong to this class detected by the pattern: ALL.
- -Other sequence(s) detected in Swiss-Prot: 38.
- -Note: Drosophila vitellogenins are also related to lipases [5], but they have lost their active site serine.
- -Last update: November 1997 / Pattern and text revised.
- [1] Chapus C., Rovery M., Sarda L., Verger R. Biochimie 70:1223-1234(1988).
- [2] Persson B., Bengtsson-Olivecrona G., Enerback S., Olivecrona T., Joernvall H. Eur. J. Biochem. 179:39-45(1989).
- [3] Blow D.
 - Nature 343:694-695(1990).
- [4] McLean J., Fielding C., Drayna D., Dieplinger H., Baer B., Kohr W., Henzel W., Lawn R. Proc. Natl. Acad. Sci. U.S.A. 83:2335-2339(1986).
- [5] Baker M.E. Biochem. J. 255:1057-1060(1988).

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